



## Rationale - Why “SmallSats”?

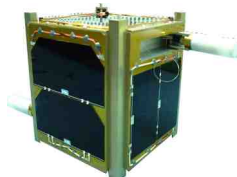


- **SmallSats are ever more capable:** *Miniature/micro/nano technology* advances
  - fabrication; materials; optics; sensors; actuators; fluidics; MEMS; electronics; communications; instrumentation; data handling & storage
  - *Power* generation & storage density up; power needs down
- **Access to space** *Low-cost launch accommodations as secondary payloads*
  - military, gov't., commercial; US, Russia, Europe, Canada, India
  - **Multiple flights possible** - test, learn, iterate
  - *Low cost* brings new participants: developing nations, universities, high schools
- **Excellent education vehicle:** Significant academic participation worldwide
- **Autonomous operations:** Less reliance on human-tended experiments
  - Eliminates possible bottleneck; can increase reliability
- **Technology migration:** ISS; other “free-flyer” satellite platforms (low Earth orbit, geostationary orbit, libration points); landers/orbiters for moon, Mars, other planets

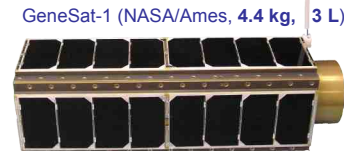


Cal Poly

“1U”  
CubeSats



PACE: National Cheng Kung University, Taiwan



GeneSat-1 (NASA/Ames, 4.4 kg, 3 L)

Launched: Dec. 2006  
Re-entry/disintegration: ~ Aug. 2010

## Astrobiology & Space Biology



**Astrobiology:** origin, evolution, distribution, & future of life in the universe

- Understand details & distribution of prebiotic chemistry -- chemical building blocks of life
- Study potential for life to adapt/survive in extraterrestrial environments
- Search for (signs of) extant or extinct non-terrestrial life
- Find habitable environments in our solar system & beyond
- **Why: fundamental understanding of life & the universe**



**Fundamental Space Biology:** effects of the space environment on terrestrial life

- Reduced (micro) gravity effects
  - Mammals: fluid distribution, musculoskeletal loading  $\Rightarrow$  immune stress, bone density decrease, muscle atrophy, slowed wound healing
  - Cells, microorganisms in culture: nutrient and waste transport
- Radiation effects: damage from (high-energy) ionizing radiation
  - Greater outside Earth's magnetosphere, ~70,000 km
  - DNA damage: strand breaks, cell death, mutations
  - Cell membrane, protein, & oxidative damage
- Bio/chemical effects of extraterrestrial environments: lunar dust
- Synergies of combined  $\mu$ gravity & radiation effects possible
- **Why: human space travel, moon/planetary habitation; insights & therapies for human disease, aging, radiation effects**

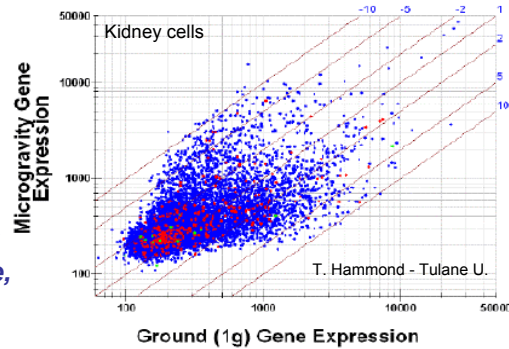


## GeneSat: Why Genetics in Space?

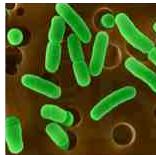


- **Current NASA mission is human exploration of the solar system**
  - Must understand adverse effects at the most fundamental levels of biology if therapies (countermeasures) are to be developed
  - *In-situ* experiments produce immediate, more accurate info. (vs. sample return)
  - **Modern genetic research is compatible with small, autonomous payloads**
- **Deleterious effects of space travel are relevant to health on Earth**
  - Osteoporosis
  - Muscle atrophy
  - Immune efficiency degradation
  - Radiation damage
  - ***Some biological effects are accelerated in space: unique insights into their mechanisms are possible, leading to therapies***

### Microgravity (Space Shuttle)

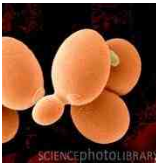


## Model Organisms



### *E. coli* (GeneSat)

- ~1 x 2  $\mu$ m bacteria
- nutrient-deprived dormancy (4 – 37  $^{\circ}$ C) until stable orbit
- GFP fusions to track expression of key genes
- diffuse fluorescence assay of GFP levels
- optical density measurement for population estimate



### *S. cerevisiae* (PharmaSat)

- ~ 5  $\mu$ m fungi, spores
- cold water stasis until stable orbit
- optical density (light scattering) tracks population density
- redox indicator dye tracks metabolic activity



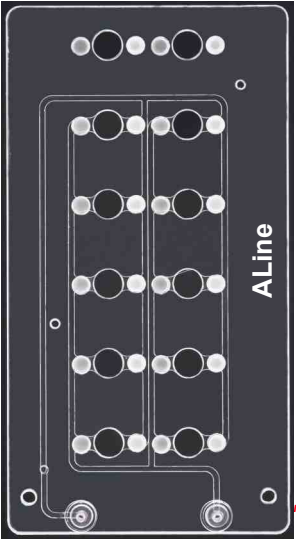
### *B. Subtilis* & *Halorubrum chaoviatoris* (O/OREOS Sat)



### *C. elegans* ( $\mu$ Satellite *in-situ* Technologies Imager)

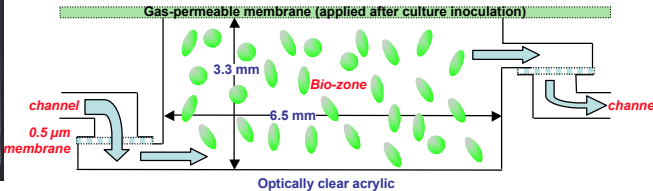
- mature adults: ~50  $\mu$ m x 1 mm
- dauer state dormancy (L3) until stable orbit
- green/red fluorescent protein fusions: gene expression
- imaging fluorescence analysis, 2 colors, ~ 9  $\mu$ m resolution

<b>GeneSat Payload System</b> <b>Fluidic card</b>	Date: 11-June-07 David Oswell, Tony Ricco, Chris Storment, Matthew Piccini, Leanna Levine (Aline)
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**ALine**

- 12-well culture-and-analysis plate
  - 10 assay wells, 2 control/standard wells
  - 110  $\mu\text{L}$ /well  $\Rightarrow$  1.1 mL total on-card volume
- Reservoir capacity  $\sim$ 15 mL
- Membrane filter at each well inlet and outlet
- Loaded pre-launch with *E. coli* in stasis medium
- Infused upon stable (*g*, T) orbit with glucose solution to initiate growth



Gas-permeable membrane (applied after culture inoculation)

3.3 mm

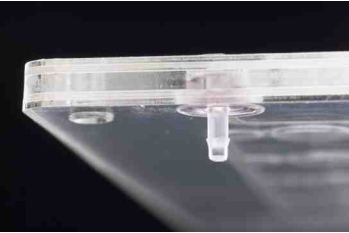
Bio-zone

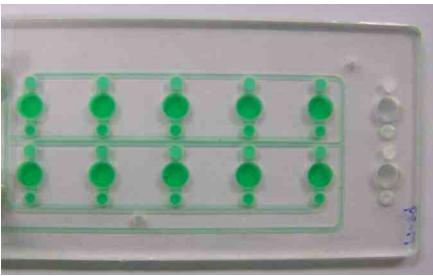
6.5 mm

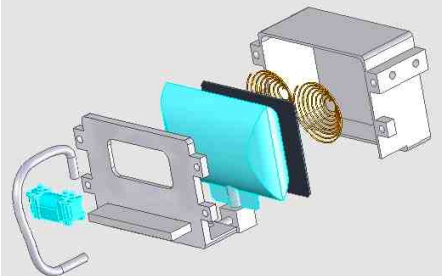
Optically clear acrylic

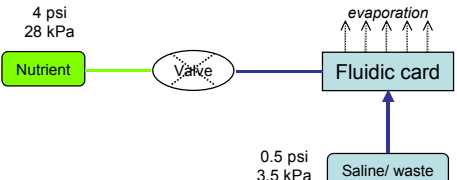
channel 0.5  $\mu\text{m}$  membrane channel

<b>GeneSat Payload System</b> <b>Fluidic system</b>	Date: 11-June-07 George Swaiss, David Oswell, Chris Storment, Matthew Piccini
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4 psi  
28 kPa

Nutrient

Valve

evaporation

Fluidic card

0.5 psi  
3.5 kPa

Saline/ waste

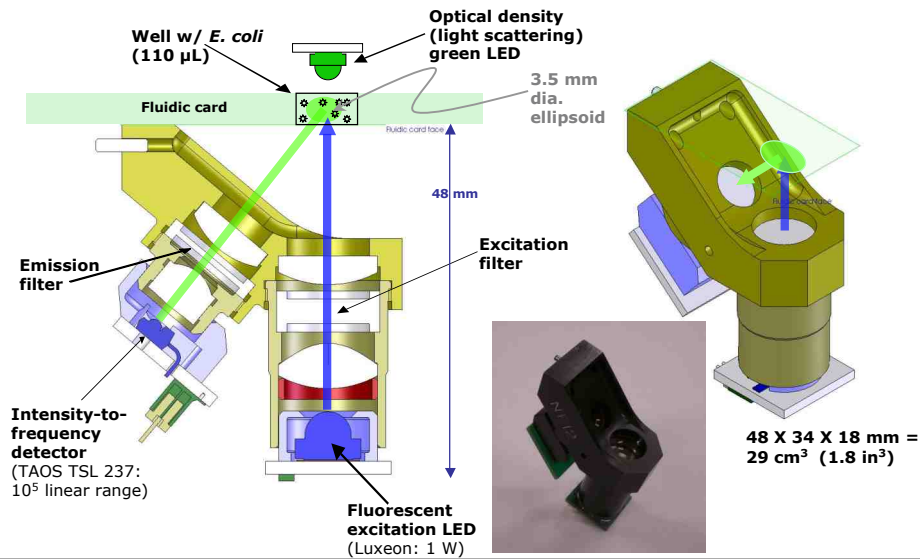
## GeneSat Payload System

Date: 11-June-07



### Optical system

Linda Timucin, Tony Ricco, Stephane Follonier, Peter Mrdjen, Bob Ricks, Optical Research Associates, Optics One



## GeneSat Payload System

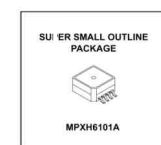
Date: 11-June-07



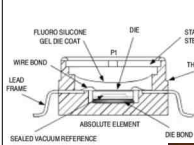
### Sensors

Chris Storment, Bob Ricks, Matthew Piccini

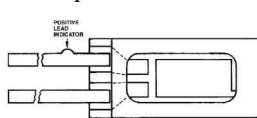
#### Pressure Sensor



#### Motorola MPXH6101A



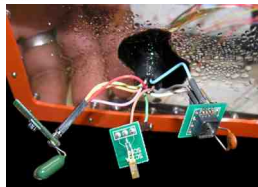
#### Temperature Sensor



#### Relative Humidity Sensor

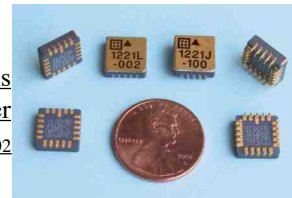


#### Radiation Sensor (PIN diode)

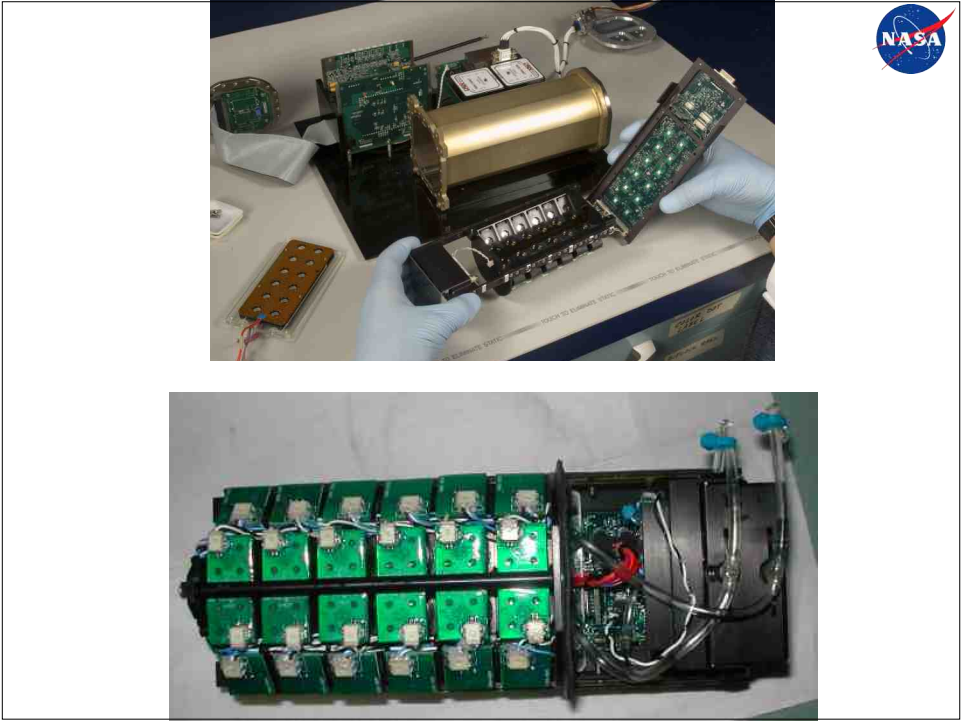


#### 3-Axis Accelerometer

Silicon Designs 1221-002





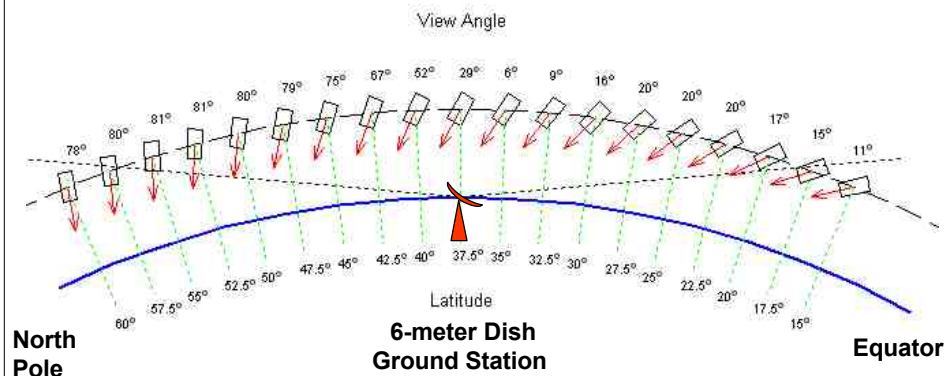


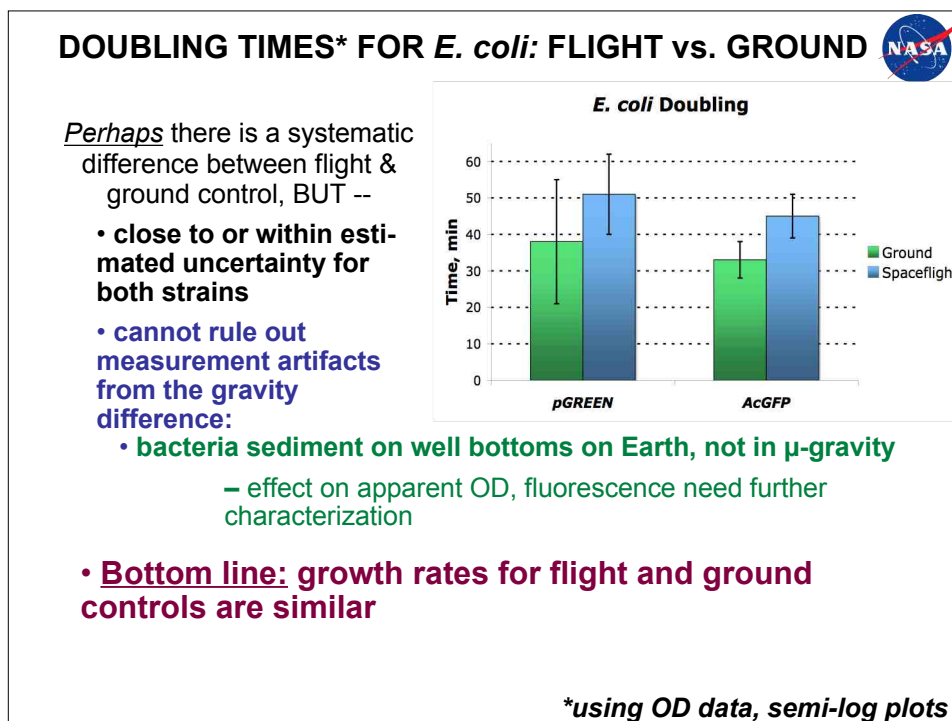
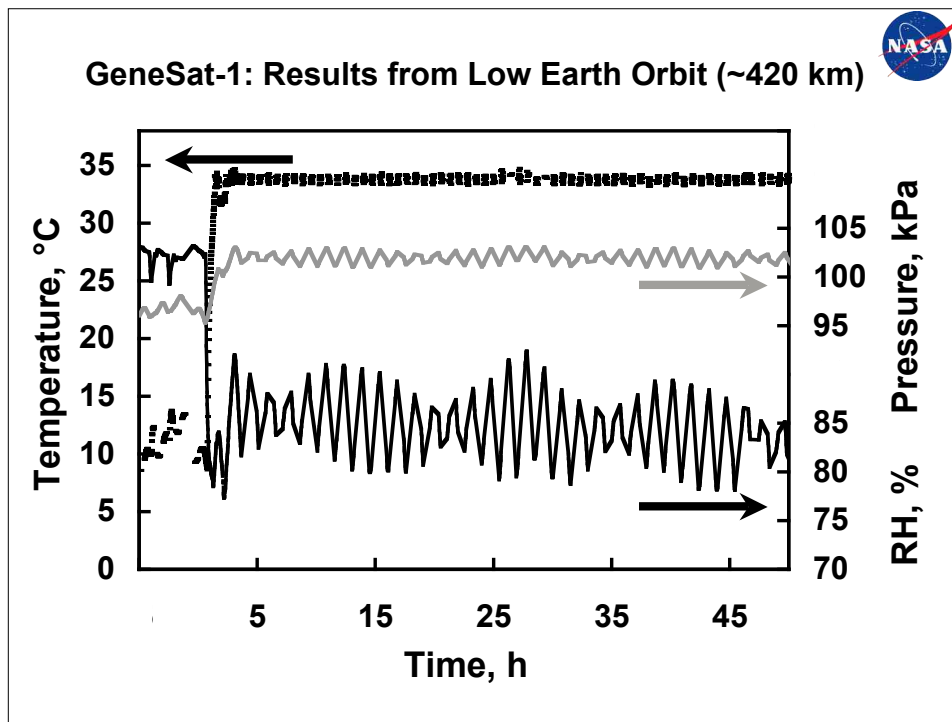
# GeneSat Launch: 16 Dec 2006



420 km, 90 min orbits; burned up on re-entry, 8-Aug-2010

## GeneSat Passive Attitude Control Points Patch Antenna Toward SRI Ground Station







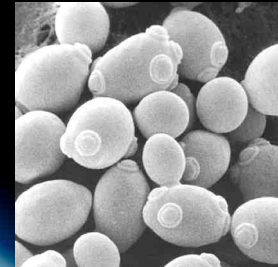


## PharmaSat: Effect of Microgravity on Yeast Susceptibility to Antifungal Drugs



Tony Ricco, Macarena Parra, John Hines, Mike McGinnis, Dave Niesel, Matthew Piccini, Linda Timucin, C. Friedericks, E. Agasid, C. Beasley, M. Henschke, C. Kitts, A. Kudlicki, E. Luzzi, D. Ly, I. Mas, M. McIntyre, R. Rasay, R. Ricks, K. Ronzano, D. Squires, J. Tucker, B. Yost

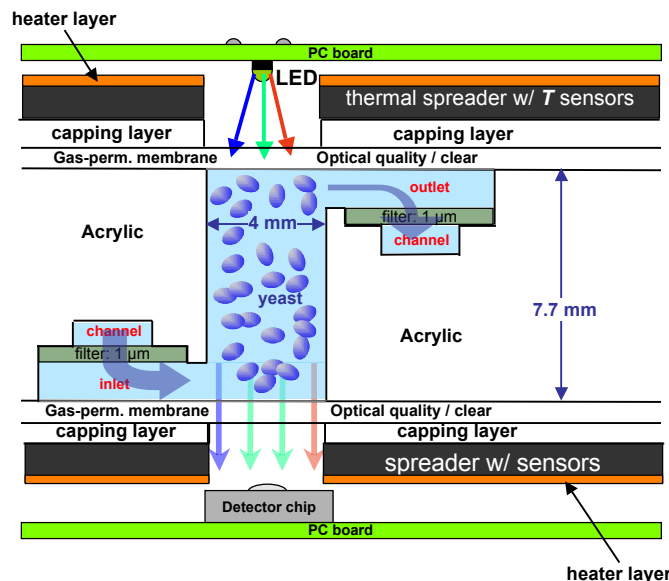
- **Grow yeast cells in multiwell fluidics card in microgravity**
  - Measure efficacy of antifungal agent to inhibit growth of fungus
    - Control + 3 concentrations of antifungal
      - 12 wells each for statistics
  - **Measure cell health & growth:**
    - Optical absorbance (turbidity, OD)
    - Viability indicator: Alamar Blue
      - Colorimetric assay: metabolic products cause blue dye → pink dye



*S. cerevisiae*

**LAUNCH: 19 May 2009**

## Fluidic/Thermal/Optical Architecture



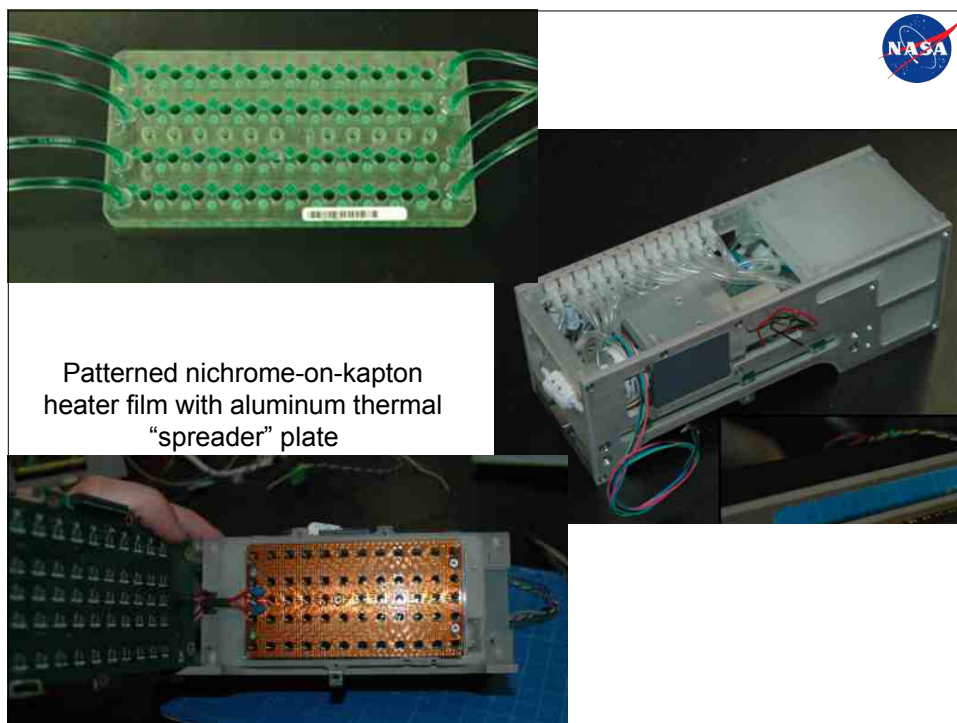
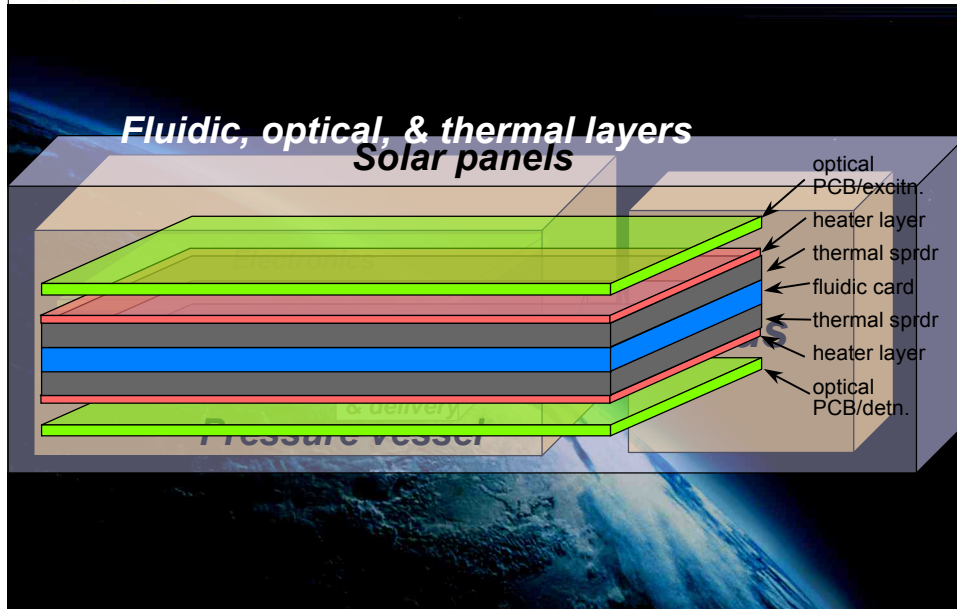
3-color LED for OD & viability: track population during growth, viability using Alamar Blue indicator dye

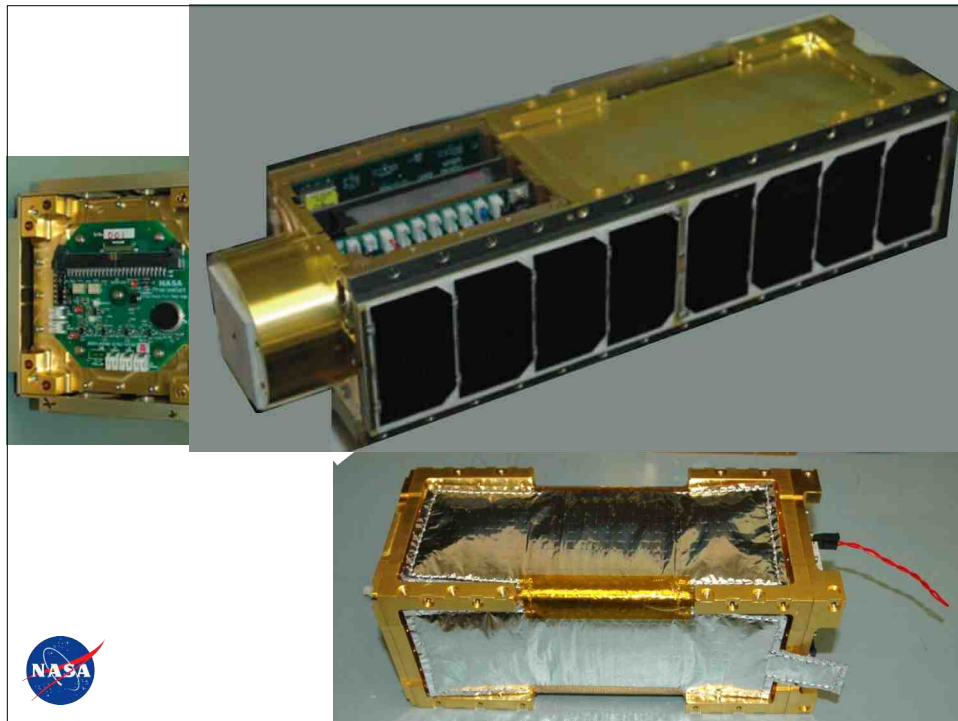
**Fluidic/  
optical/  
thermal  
cross-  
section**

Detector for OD and viability measurement using 3-color absorbance



## PharmaSat Technology Architecture - 2





## PharmaSat Launch: ***19 May 2009***



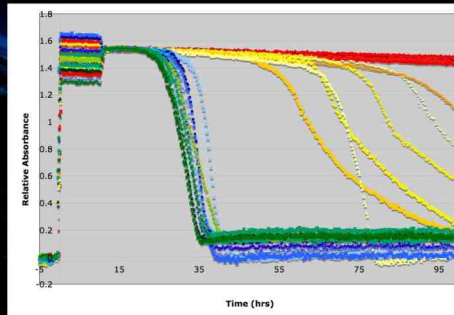
**420 km, 90 min orbits; still up there (for a while...)**



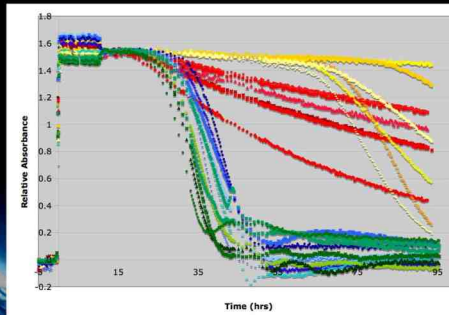
## PharmaSat Data (*red channel*)



### Ground



### Spaceflight



■ Hi Antifungal

■ MIC\* Antifungal

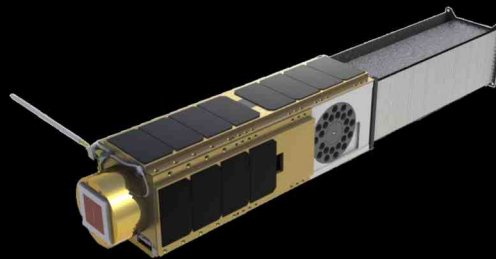
■ Lo Antifungal

■ Control

\*MIC = minimum inhibitory conc.

- Antifungal effect is clear in both Spaceflight and Ground
- Response for High Antifungal consistent with metabolism being less suppressed in  $\mu$ gravity than on Earth... but cell division remains suppressed

## Organism/ORganics Exposure to Orbital Stresses: The O/OREOS NanoSatellite



A. J. Ricco, D. Squires, J. W. Hines, P. Ehrenfreund,<sup>1</sup>  
R. Mancinelli<sup>2</sup>, A. Mattioda, W. Nicholson<sup>3</sup>, R. Quinn<sup>2</sup>, O. Santos

Science support: N. Bramall, J. Chittenden, K. Bryson, A. Cook, M. Parra, D. Ly

Development by the NASA-Ames  
Small Spacecraft Payloads & Technologies Team

NASA Ames Research Center

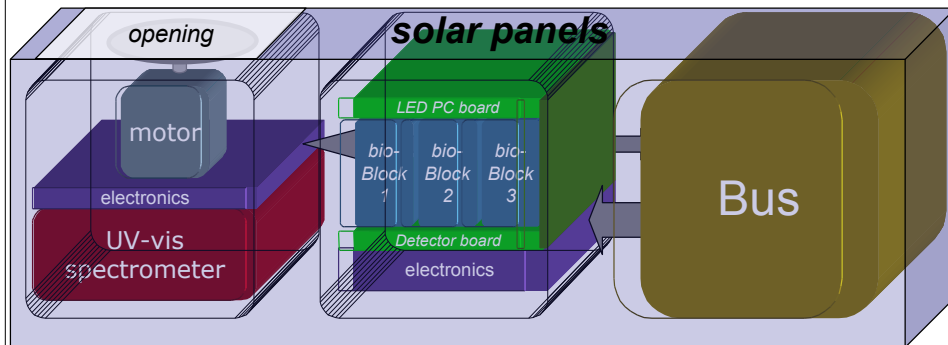
<sup>1</sup> George Washington University

<sup>2</sup> SETI Institute

<sup>3</sup> University of Florida/Kennedy Space Center

## O/OREOS Dual-Payload Technology Architecture

*Each P/L experiment-plus-instrument contained in a single 10-cm cube*



### Organics Payload

- 4 organics
- 4 environments
- UV-vis. spectrometer

### Biology Payload

- 2 biological specimens
- 3 growth initiation times
- optical measurements

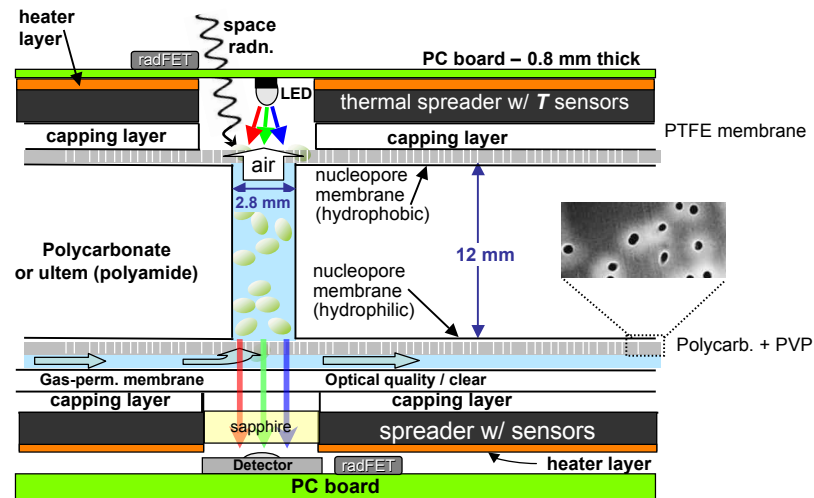
## Payload 1: Space Environment Survivability of Live Organisms (SESLO)

- Two organisms, wildtype & mutant, exposed to  $\mu$ gravity & space radiation
  - $< 10^{-3}$  x Earth gravity
  - 2 – 20 Gy total dose over 6 mo in 650 km orbit
    - 0.1 Gy is galactic cosmic radiation
- Dry organisms on  $\mu$ well walls pre integration
- Rehydrate & feed 6  $\mu$ wells / organism:  $t = 2$  wk, 3 mo, 6 mo
- Grow @  $35 - 37 \pm 2$  °C for 1 – 17 days
- Measure RGB transmittance @ 615, 525, 470 nm
  - track culture population via calculated optical density (both organisms)
  - track metabolic activity via Alamar Blue
- Sensors:  $T$ ,  $p$ ,  $RH$ ,  $rad$  (integrated dose),  $\mu$ grav
  - » temperature (6 sensors per 12-well bioblock)
  - » pressure, relative humidity (1 sensor each)
  - » radiation total dose @ both ends of wells (2 radFETs)
  - » microgravity levels calc'd. from solar panel currents

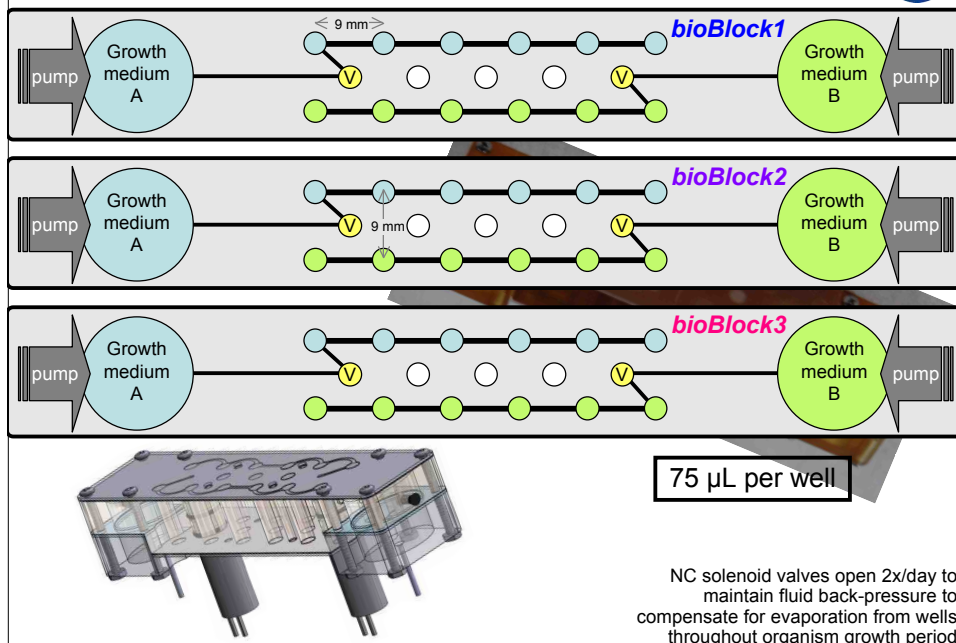


## SESLO (bio) Fluidic/Thermal/Optical Architecture

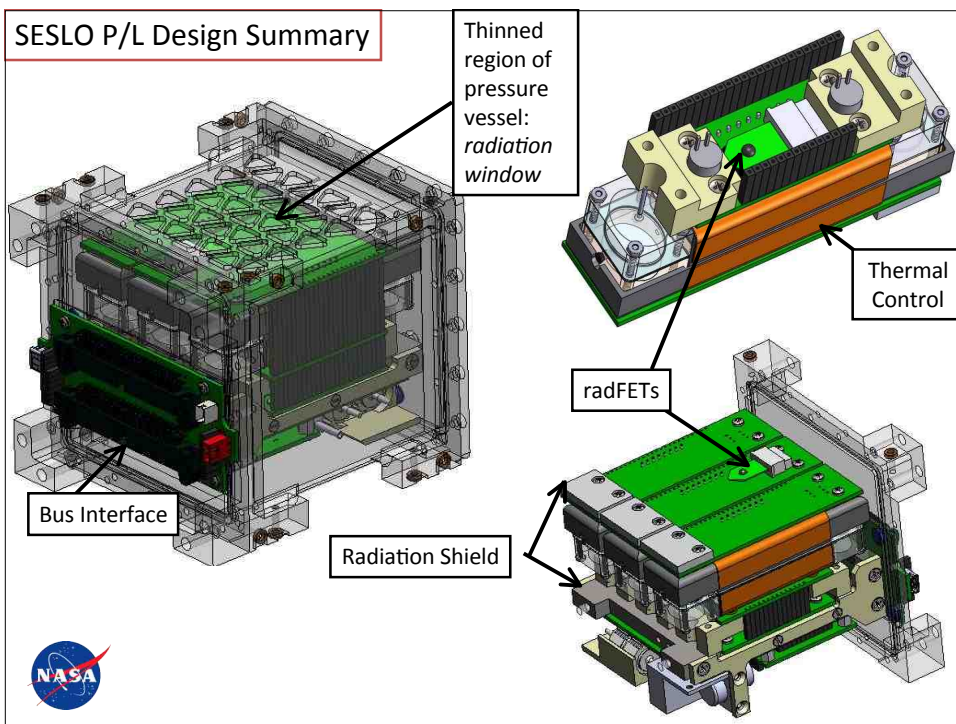
### Fluidic / optical / thermal cross-section



## SESLO Integrated Fluidic System: 3 independent bioBlocks

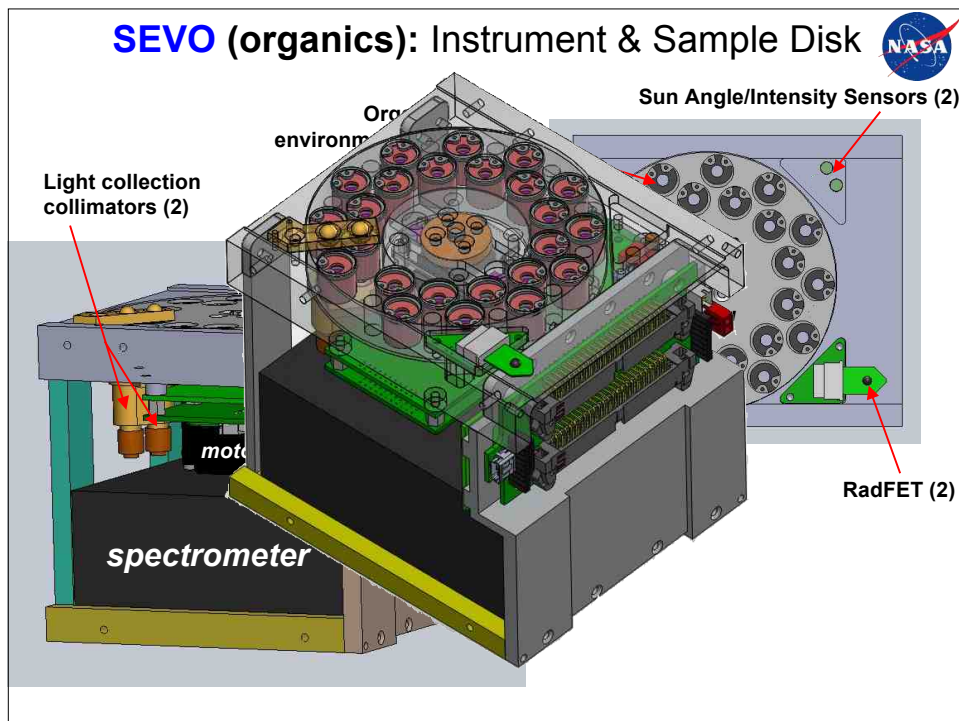
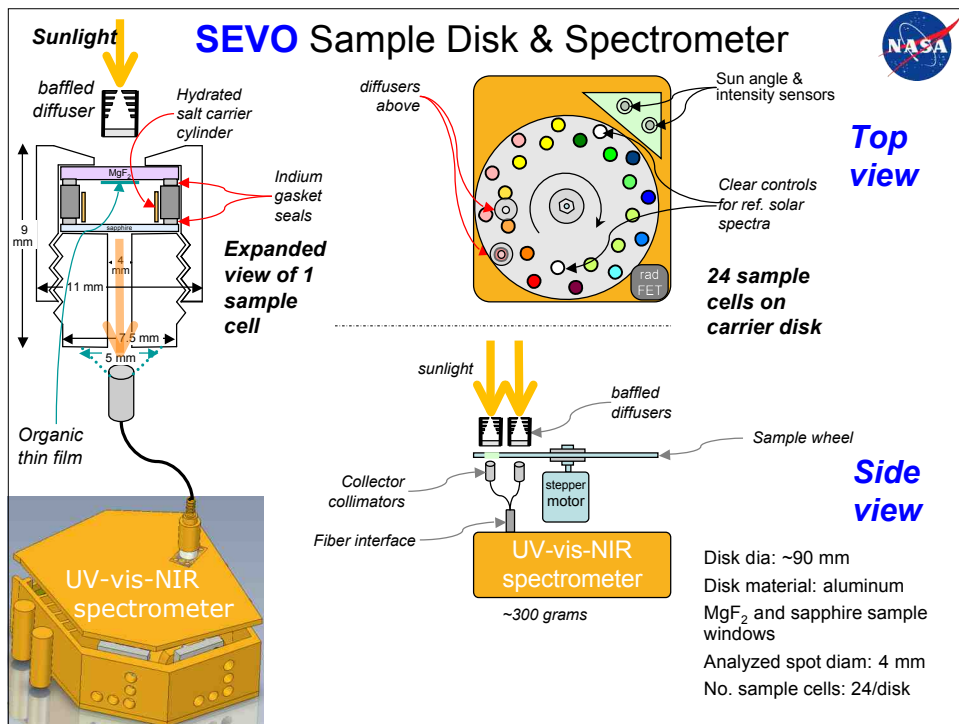






**Payload 2: Space Environment Viability of Organics (SEVO)**

- **6-mo exposure** of organics to space
  - amino acid; polyaromatic hydrocarbon; quinone; porphyrin
  - 4 self-contained sample micro-environments: **interstellar** (vacuum), **lunar**, **martian**, **brine** (humidity only)
  - ionizing radiation: ~ 50 Gy = 5 krad
  - solar exposure 20 – 35% of total time, ~120 - 2500 nm
- Measure **UV-vis-NIR spectrum daily** over 6-mo mission
  - for all 18 organic films & 6 references, 200 - 1000 nm
  - sun is light source for spectroscopy
  - solar spectral features: internal wavelength calibration
- Sensors:  $T_{sample}$ ,  $T_{spec}$ ,  $I_{solar}$ ,  $\theta_{solar}$ ,  $rad$  (integrated dose)
  - temperature: multiple sensors, spectrometer & sample wheel
  - sun angle: acquire spectra only with appropriate illumination
  - solar intensity: calibration with each sample spectrum
  - radiation total dose: near cells



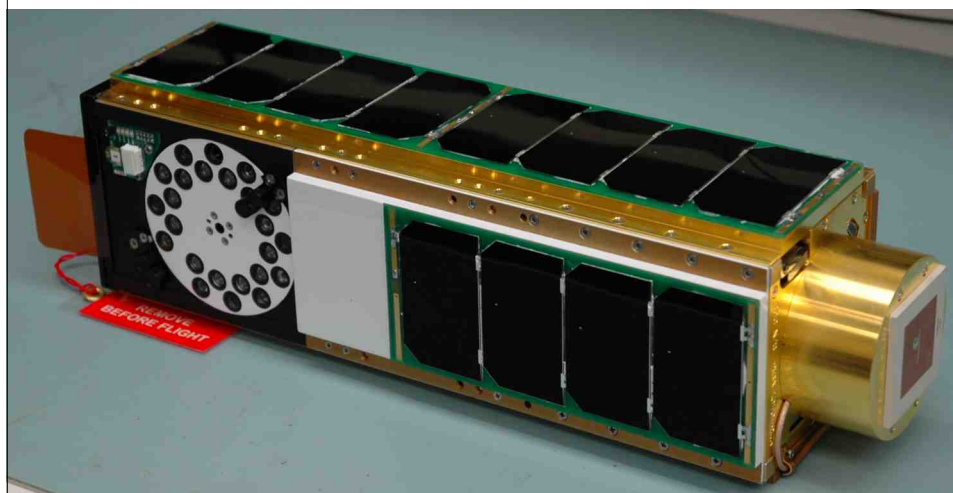
## O/OREOS Flight Prototype



SEVO

SESLO

## O/OREOS Nanosatellite



*Launch: November 19, 2010 (Kodiak, Alaska)*

## Conclusions



**The tools of bio- and micro-technology combined with automation & integration enable a range of biology experiments in small space platforms**

- **Small satellites enable more experiments: space access, low cost**
- **Fundamental biological effects/phenomena explored in a unique environment**
  - Human health & safety
  - Origin, evolution, distribution, & future of life in the universe (astrobiology)
  - Unique zero-shear-rate suspensions of cells & microorganisms
- **Relevance to terrestrial medicine & pharma development**
  - Accelerated test platform: osteoporosis, muscle atrophy, immune impairment, radiation effects
  - Novel conditions impact microorganism function including metabolic processes, secreted proteins
  - Spaceflight environment increases the virulence of some pathogens
  - Growth of ultra-low-defect-density protein crystals